

Public
gather
collecti
Data is

1 hour per response, including the time for reviewing instructions, searching existing data sources, collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Avenue, Washington, DC 20540, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

1. AG

6/28/94

3. REPORT TYPE AND DATES COVERED
Technical Report4. TITLE AND SUBTITLE
Synthesis of Polyphenylenes and
Polynaphthalenes by Thermolysis of Eneidyne and
Dialkynylbenzenes5. FUNDING NUMBERS
G- N00014-89-J3062
R&T 3132084
K. Wynne

6. AUTHOR(S)

Jens A. John and James M. Tour

7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)
Department of Chemistry and Biochemistry
University of South Carolina
Columbia, SC 29208

94-20877

9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES)
Department of the Navy
Office of Naval Research
800 North Quincy Street
Arlington, VA 22217-500010. SPONSORING/MONITORING
AGENCY REPORT NUMBER

30

11. SUPPLEMENTARY NOTES

J. Am. Chem. Soc., 1994, 116, 5011.

12a. DISTRIBUTION/AVAILABILITY STATEMENT

Reproduction in whole or in part is permitted for any
purpose of the United States Government. This document has
been approved for public release and sale; its distribution
is unlimited.

12b. DISTRIBUTION CODE

13. ABSTRACT (Maximum 200 words)

Described are the syntheses of substituted eneidyne and dialkynylbenzenes using Pd- or Pd/Cu-catalyzed cross coupling procedures. The products were then thermalized, generally in benzene, to afford the corresponding poly(p-phenylene)s and poly(1,4-naphthalene)s. Thirteen examples are provided that show the scope of the polymerization process based upon substituent patterns and cyclization moieties. The superb thermal resiliency of the newly derived polymers is demonstrated using thermogravimetric analysis. The polymer structure was generally confirmed using IR data correlations to small molecules that resembled the polymer's repeat unit structure. Radical trapping of dimeric intermediates, that were analyzed by GCMS, further substantiated the proposed mechanistic route. The step-growth polymerization pattern was determined by monitoring the degree of monomer consumption versus the polymer molecular weight.

DTIC QUALITY INSPECTED 5

14. SUBJECT TERMS

15. NUMBER OF PAGES

16. PRICE CODE

17. SECURITY CLASSIFICATION
OF REPORT
Unclassified18. SECURITY CLASSIFICATION
OF THIS PAGE
Unclassified19. SECURITY CLASSIFICATION
OF ABSTRACT
Unclassified20. LIMITATION OF ABSTRACT
UL

94 7 7 077

OFFICE OF NAVAL RESEARCH

GRANT : N00014-89-J-3062

R&T Code 3132084

Technical Report No. 30

Synthesis of Polyphenylenes and Polynaphthalenes by Thermolysis of Eneidyne and
Dialkynylbenzenes

by

Jens A. John and James M. Tour

Accepted for Publication in

Journal of the American Chemical Society

Department of Chemistry and Biochemistry
University of South Carolina
Columbia, SC 29208

June 28, 1994

Reproduction in whole, or in part, is permitted for any purpose of the United States Government

This document has been approved for public release and sale; its distribution is unlimited.

Accession For	
NTIS CRA&I	<input checked="" type="checkbox"/>
DTIC TAB	<input type="checkbox"/>
Unannounced	<input type="checkbox"/>
Justification _____	
By _____	
Distribution /	
Availability Codes	
Dist	Avail and/or Special
A-1	

**Synthesis of Polyphenylenes and Polynaphthalenes by Thermolysis of Eneidyne
and Dialkynylbenzenes**

Jens A. John and James M. Tour*

Department of Chemistry and Biochemistry

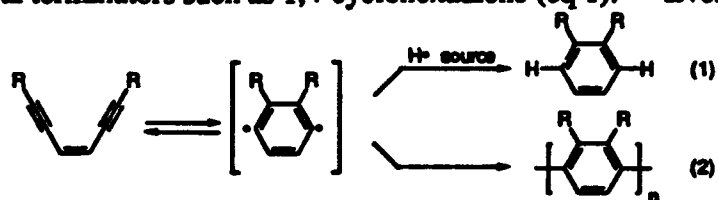
University of South Carolina

Columbia, South Carolina 29208

Abstract

Described are the syntheses of substituted eneidyne and dialkynylbenzenes using Pd- or Pd/Cu-catalyzed cross coupling procedures. The products were then thermalized, generally in benzene, to afford the corresponding poly(*p*-phenylene)s and poly(1,4-naphthalene)s. Thirteen examples are provided that show the scope of the polymerization process based upon substituent patterns and cyclization moieties. The superb thermal resiliency of the newly derived polymers is demonstrated using thermogravimetric analysis. The polymer structure was generally confirmed using IR data correlations to small molecules that resembled the polymer's repeat unit structure. Radical trapping of dimeric intermediates, that were analyzed by GCMS, further substantiated the proposed mechanistic route. The step-growth polymerization pattern was determined by monitoring the degree of monomer consumption versus the polymer molecular weight.

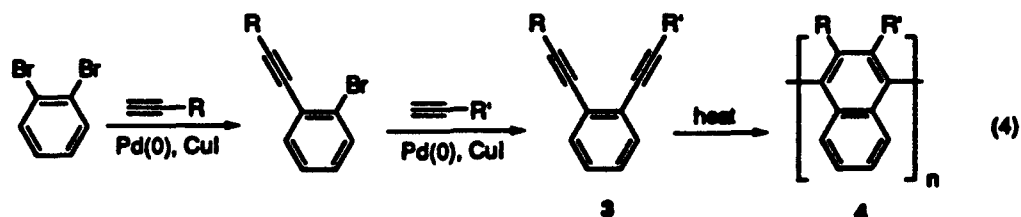
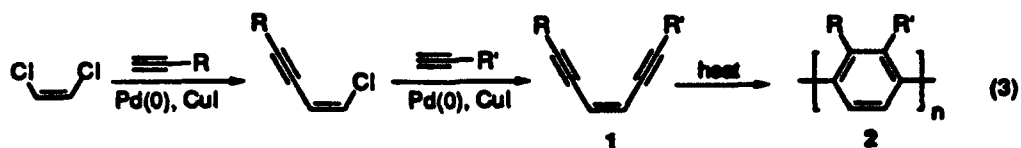
Bergman's study on the thermal cycloaromatization of enediynes led to the suggestion of a benzene 1,4-diradical intermediate.¹ Numerous synthetic and mechanistic studies were later performed to further investigate this reaction and a variety of enediynes have been thermalized in the presence of radical terminators such as 1,4-cyclohexadiene (eq 1).^{1,2} Even though large



excesses of radical terminators were employed, the yield of the substituted benzenes were often moderate at best. The fact that the yields were diminutive kindled our interest because it suggested to us that polymerization may indeed be a preferred pathway and we could regard the intermediate benzene 1,4-diradical as a building block for substituted polyphenylenes. As suspected, deletion of the radical trapping source from the reaction mixtures allowed for the formation of substituted polyphenylenes (eq 2).³

Substituted polyphenylenes have been shown to exhibit superb thermal and chemical resilience, interesting semiconducting properties upon doping, and applications in light emitting diodes.^{4,5} Therefore, this route to polyphenylenes and their derivatives is most attractive since (1) it requires no exogenous chemical catalysts or reagents for the polymerization, (2) no heteroatomic coupling sites such as halogens are not necessary and all atoms present in the monomer are also present in the polymer, and (3) the monomers can be rapidly prepared with a variety of substitution patterns to often afford soluble polyphenylene derivatives.

We could rapidly prepare numerous enediynes and dialkynylbenzenes by the Pd/Cu coupling protocol or an alkynylzinc chloride/Pd coupling.^{6,7} Subsequent heating of the enediynes or dialkynylaromatics lead to polymerization products (eqs 3 and 4). The results of the polymerization are summarized in Table I. We found that highest molecular weights could be obtained using benzene as a solvent and heating the monomers in thick-walled screw cap glass tubes⁸ at 50-160°C. In a few cases, when the polymerizations required higher temperatures, they were conducted on neat materials.



In general, monomers that had both R and R' = H, afforded insoluble polymers. Monomers that had either R or R' = H, readily polymerized to afford soluble polymers. In the cases where neither R or R' = H, the polymerizations were often slower and they required higher temperatures and neat reaction conditions which likely caused polymer decomposition as flagged by the black color of the products (Table I). The yields of the polymerized material were generally 50-90% after fractional precipitation. The obtained M_n values of 1500-2500 are quite typical for *p*-polyphenylene derivatives.⁴ Note, however, that SEC-determined values of M_n versus polystyrene standards are inflated due to hydrodynamic volume differences of rigid rod polymers versus the randomly coiled polystyrene standards.

The superb thermal stabilities are apparent in that, in many cases, the unannealed polymers exhibited 10% weight loss only after exceeding 400-600°C (Table I). Being unannealed, some of the early weight loss could be attributed to occluded solvent, therefore, these values represent a lower limit for thermal degradation.

The IR spectra are most informative in confirming the structure of the proposed polymers.⁹ For the monosubstituted polyphenylenes (Table I, entries 1-7), the polymer repeat unit possesses a 1,2,4-substituted phenyl pattern. The most diagnostic bands at 935-810 cm^{-1} are assigned to the bending of a lone hydrogen at the 3-position and the band at 880-795 cm^{-1} arises from the bending of two the adjacent hydrogens at the 5- and 6-positions. Assignment of the disubstituted polyphenylenes can best be described using the bis(trimethylsilyl) derivative (Table I, entry 9). The shoulder above 3000 cm^{-1} comes from the C-H stretching vibration. The band at 838 cm^{-1}

results from out-of-plane C-H bending of two adjacent hydrogen atoms bound to a phenyl ring. Analysis of the 1,2-dialkynylbenzene-derived polymers argued in favor of the 1,4-disubstituted naphthalene structure (eq 4). For example, for the simplest case where $R = R' = H$ (Table I, entry 10), bands at 1595, 875 (two adjacent H) and 754 (four adjacent H) agreed with the literature values for 1,4-disubstituted naphthalenes.⁹

The mechanistic aspects of the polymerization process are intriguing.¹⁰ Two general mechanisms are considered. First, a benzene 1,4-diradical may be attacking unreacted enediyne to generate a new biphenyl 4,4'-diradical intermediate. Continued reaction of this growing chain with enediyne should result in a chain growth-like polymerization process in that the molecular weight of the polymer would increase rapidly even at low monomer conversions. Conversely, the benzene 1,4-diradical may predominantly couple with other radicals in solution to eventually form the polymer. In that case, the process should be step growth-like where the polymer molecular weight would initially increase slowly, but then increase rapidly when monomer conversion was nearly complete. By monitoring consumption of **1** ($R = H$, $R' = Ph$) (using capillary GC with terphenyl as an internal standard) and molecular weight (M_n by SEC), it was clear that the molecular weight increased slowly ($M_n = 600$ at 10% monomer consumption and 810 at 90% monomer consumption) until the monomer was nearly consumed (>97%), whereupon the molecular weight increased exceedingly rapidly. Therefore, the polymerization indeed resembles a step growth process. It should be noted, however, a chain growth mechanism can not be ruled out in that the propagating species may be unstable giving rapid chain transfer and/or termination. Also, addition of small amounts of radical initiators did not increase the polymerization efficiency.

As a further conformation of the radical coupling process described here, we carried out the thermalization process for two of the monomers in the presence of some of the radical trapping agent 1,4-cyclohexadiene. This was done with the hope of obtaining some small, characterizable dimeric species. Indeed, when **1** ($R = H$, $R' = Ph$) was heated in benzene in the presence 1,4-cyclohexadiene (33 mol %) at 150°C, we observed the formation of polymeric material as well as several low molecular weight species detected by GC/MS analysis. The volatile material consisted

of unreacted starting material, biphenyl (154 amu) and the desired dimeric adduct quaterphenyl (306 amu). Likewise, when **3** (R and R' = H) was heated in benzene in the presence 1,4-cyclohexadiene (21 mol %) at 150°C, polymer was formed as well as several volatile species. The volatile material consisted of unreacted starting material, naphthalene (128 amu), 1,4-cyclohexadienyl addition/aromatization products, and the desired dimer 1,1'-binaphthalene (254 amu). The isotopic pattern and fragmentation pattern matched identically with the data base-derived 1,1'-binaphthalene spectrum.

Finally, it has been shown that *E*-enediynes do not undergo the Bergman cyclization.¹ We therefore prepared *E*-1-phenylhexa-3-ene-1,5-diyne. Heating this monomer to 150°C for 2 d did indeed form a polymer, however, FTIR analysis verified that it was not a polyphenylene-derived material in that the two bands at 935-810 cm⁻¹ and 880-795 cm⁻¹ were missing.

Acknowledgments. We are grateful for support from the Office of Naval Research and the National Science Foundation (EHR-91-08772, DMR-9158315), and generous industrial contributors to the NSF Presidential Young Investigator Award (1991-96): Hercules, IBM, Ethyl, Shell, Eli Lilly, Polaroid, and Farchan Corporations. We also thank Molecular Design Ltd. for the use of their synthetic data base.

References and Notes

(1) Lockhart, T. P.; Comita, P. B.; Bergman, R. G. *J. Am. Chem. Soc.* **1981**, *103*, 4082. Lockhart, T. P.; Bergman, R. G. *J. Am. Chem. Soc.* **1981**, *103*, 4091. Bergman, R. G. *Acc. Chem. Res.* **1973**, *6*, 25. Jones, R. R.; Bergman, R. G. *J. Am. Chem. Soc.* **1972**, *94*, 660.

(2) Lee, M. D.; Dunne, T. S.; Chang, C. C.; Ellestad, G. A.; Siegel, M. M.; Morton, G. O.; McGahren, W. J.; Borders, D. B. *J. Am. Chem. Soc.* **1987**, *109*, 3466. Golik, J.; Dubay, G.; Groenewold, G.; Kawaguchi, H.; Konishi, M.; Krishnan, B.; Ohkuma, H.; Saitoh, K.; Doyle, T. W. *J. Am. Chem. Soc.* **1987**, *109*, 3462. Beau, J. M.; Crevisy, C. *Tetrahedron Lett.*

1991, 32, 3171. Wender, P. A.; McKinney, J. A.; Mukai, C. *J. Am. Chem. Soc.* 1990, 112, 5369. Nicolaou, K. C.; Dai, W.-M. *Angew. Chem. Int. Ed. Engl.* 1991, 30, 1387.

(3) (a) Recently, Bergman reported a double cycloaromatization based on a conjugated dienetriyne. See: Bharucha, K. N.; Marsh, R. M.; Minto, R. E.; Bergman, R. G. *J. Am. Chem. Soc.* 1992, 114, 3120. (b) Likewise, recent tandem cyclizations have been accomplished by Grissom. See: Grissom, J. W.; Calkins, T. L.; McMillen, H. A. *J. Org. Chem.* 1993, 58, 6556. Grissom, J. W.; Klingberg, D. *J. Org. Chem.* 1993, 58, 6559. (c) Swager and Grubbs used a similar cascade sequence to cycloaromatize a poly(arylene-ethynylene). See: Zhou, Q.; Swager, T. M. *Polym. Prepr. (Am. Chem. Soc., Div. Polym. Chem.)* 1993, 34, 193. Grubbs, R. H.; Kratz, D. *Chem. Ber.* 1993, 126, 149.

(4) For some reviews on polyphenylene, see: (a) Kovacic, P.; Jones, M. B. *Chem. Rev.* 1987, 87, 357. (b) Noren, G. K.; Stille, J. K. *Macromol. Rev.* 1971, 5, 385. (c) Elsenbaumer, R. L.; Shacklette, L. W. In *Handbook of Conducting Polymers*; Skotheim, T. A., Ed.; Dekker: New York, 1986. (d) Tourillon, G. in ref 5c. (e) Baughman, R. H.; Brédas, J. L.; Chance, R. R.; Elsenbaumer, R. L.; Shacklette, L. W. *Chem. Rev.* 1982, 82, 209.

(5) For use in light emitting diodes, see: Grem, G.; Günther, L.; Ullrich, B.; Leising, G. *Adv. Mater.* 1992, 4, 36.

(6) (a) Sonogashira, K.; Tohda, Y.; Hagihara, N.; *Tetrahedron Lett.* 1975, 4467. Stephans, R. D.; Castro, C. E. *J. Org. Chem.* 1963, 28, 3313. Suffert, J.; Ziessel, R. *Tetrahedron Lett.* 1991, 32, 757. (b) The terminal alkyne units on the monomers were affixed to the alkenyl or aryl halides using trimethylsilylacetylene followed by TMS removal with $K_2CO_3/MeOH$.

(7) In most cases involving alkyne coupling to 1,2-dibromobenzene, we could obtain higher yields using the Pd(0)-catalyzed coupling of the alkynylzinc chloride to the arylbromide. See: Negishi, E.; Takahashi, T.; Baba, S.; Van Horn, D. E.; Okukado, N. *J. Am. Chem. Soc.* 1987, 109, 2393.

(8) CAUTION: The threaded glass tubes were obtained from Ace Glass Inc., product number 8648. Though we experienced no explosions in more than 50 reaction runs studied, thermalization processes should be carried out behind protective shielding.

(9) (a) Bellamy, L. J. *The Infra-red Spectra of Complex Molecules*, Wiley: New York, 1975. (b) Lin-Vien, D.; Colthup, N. B.; Fateley, W. G.; Grasselli, J. G. *The Handbook of Infrared and Raman Characteristic Frequencies of Organic Molecules*, Academic: New York 1991. (c) Mayer, R.; Kleinert, H.; Richter, S.; Gewald, K. *Angew. Chem., Internat. Ed. Engl.* 1962, 1, 115. (d) *The Aldrich Library of FT-IR Spectra, Edition I*, Pouchert, C. J., Ed., Aldrich Chemical Company: Milwaukee, 1985.

(10) Grissom, J. W.; Calkins, T. L. *J. Org. Chem.* 1993, 58, 5422.

Table I. Thermal Polymerization of Eneidyne and Dialkynylarenes to Afford Polyaromatics.^a

Entry	Monomer	R	R'	Temp ^b	M _w ^{c,d}	M _n ^{d,e}	solubility ^f	T _{10%} ^{g,h}	T _{50%} ^{h,i}	color ^j
1	1	H	<i>n</i> -C ₆ H ₁₃	120	3930	1570	soluble	287	472	brown
2	1	H	Ph	150	30500 ^k	9500	soluble	392	>900	tan
3	1	H	(<i>p</i> -C ₆ H ₄)-Ph	95	10600	4100	soluble	438	>900	tan
4	1	H	(<i>o</i> -C ₆ H ₄)-Ph	50-90	3200	2000	soluble	432	791	tan
5	1	H	(<i>p</i> -C ₆ H ₄)-CMe ₃	50-95	6500	2600	soluble	428	898	brown
6	1	H	(1-naphthyl)	115	5500	1450	soluble	416	>900	tan
7	1	H	(9-anthryl)	120	9400	800	partial soluble ^l	351	>900	orange ^m
8	1	<i>n</i> -C ₆ H ₁₃	<i>n</i> -C ₆ H ₁₃	140	980	810	soluble ⁿ	178	375	green
9	1	SiMe ₃	SiMe ₃	300 ^o	2850	1040	soluble	248	490	black
10	3	H	H	140	---	---	insoluble	500	>900	brown
11	3	H	Ph	140	1200	850	soluble	380	617	tan
12	3	Ph	Ph	400 ^o	---	---	insoluble	606	>900	black
13	3	<i>n</i> -C ₆ H ₁₃	<i>n</i> -C ₆ H ₁₃	340 ^o	---	---	insoluble	507	>900	black

^aDashes signify that the value was not obtained or it was not clearly discernable during the analysis. Unless otherwise noted, all reactions were carried out in a thick-walled screw-capped tube in benzene (0.7-1.1 M) as the solvent. ^bTemperature in °C used for the polymerization. The reaction times were generally 24-48 h. ^cWeight average molecular weights. ^dDetermined by size exclusion chromatography (SEC) in THF versus polystyrene standards. ^eNumber average molecular weights. ^fSolubility in THF or CH₂Cl₂. ^gTemperature at which 10% weight loss occurred. ^hDetermined by thermogravimetric analysis (TGA) from 50 to 900°C at 10°C/min under an N₂ atmosphere. ⁱTemperature at which 50% weight loss occurred. ^jColor of neat polymer. ^kWe presently have no explanation for the very broad polydispersity. ^lSEC data obtained on the soluble portion and TGA data obtained on the insoluble portion. ^mThe monomer had an intense orange color. ⁿMaterial was predominantly unreacted monomer. ^oThe reaction was run without solvent.